

PATENT COOPERATION TREATY

PCT

INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY

(Chapter II of the Patent Cooperation Treaty) D 23 MAR 2005

(PCT Article 36 and Rule 70)

WPO PCT

Applicant's or agent's file reference 4-33162A	FOR FURTHER ACTION See Form PCT/PEA/416																	
International application No. PCT/EP2004/003806	International filing date (day/month/year) 08.04.2004	Priority date (day/month/year) 10.04.2003																
International Patent Classification (IPC) or national classification and IPC C07D243/38, A61K31/551, A61P3/00																		
Applicant NOVARTIS AG et al.																		
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Date of submission of the demand 14.10.2004		Date of completion of this report 30.03.2005																
Name and mailing address of the International preliminary examining authority:  European Patent Office D-80298 Munich Tel. +49 89 2399 - 0 Tx 523658 epmu d Fax: +49 89 2399 - 4465		Authorized Officer Telephone No. +49 89 2399- 																

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INTERNATIONAL PRELIMINARY REPORT ON PATENTABILITY
(Chapter II of the Patent Cooperation Treaty)

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**INTERNATIONAL PRELIMINARY REPORT
ON PATENTABILITY**International application No.
PCT/EP2004/003806**Box No. I Basis of the report**

1. With regard to the language, this report is based on the international application in the language in which it was filed, unless otherwise indicated under this item.
 - This report is based on translations from the original language into the following language, which is the language of a translation furnished for the purposes of:
 - international search (under Rules 12.3 and 23.1(b))
 - publication of the international application (under Rule 12.4)
 - international preliminary examination (under Rules 55.2 and/or 55.3)
2. With regard to the elements* of the international application, this report is based on (replacement sheets which have been furnished to the receiving Office in response to an invitation under Article 14 are referred to in this report as "originally filed" and are not annexed to this report):

Description, Pages

1-40 as originally filed

Claims, Numbers

1-12 as originally filed

- a sequence listing and/or any related table(s) - see Supplemental Box Relating to Sequence Listing
- 3. The amendments have resulted in the cancellation of:
 - the description, pages
 - the claims, Nos.
 - the drawings, sheets/figs
 - the sequence listing (specify):
 - any table(s) related to sequence listing (specify):
- 4. This report has been established as if (some of) the amendments annexed to this report and listed below had not been made, since they have been considered to go beyond the disclosure as filed, as indicated in the Supplemental Box (Rule 70.2(c)).
 - the description, pages
 - the claims, Nos.
 - the drawings, sheets/figs
 - the sequence listing (specify):
 - any table(s) related to sequence listing (specify):

* If item 4 applies, some or all of these sheets may be marked "superseded."

INTERNATIONAL PRELIMINARY REPORT
ON PATENTABILITY

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Box No. III Non-establishment of opinion with regard to novelty, inventive step and industrial applicability

1. The questions whether the claimed invention appears to be novel, to involve an inventive step (to be non-obvious), or to be industrially applicable have not been examined in respect of:
- the entire international application,
- claims Nos. 12
- because:
- the said international application, or the said claims Nos. 12 with respect to industrial applicability relate to the following subject matter which does not require an international preliminary examination (specify):
see separate sheet
- the description, claims or drawings (*Indicate particular elements below*) or said claims Nos. are so unclear that no meaningful opinion could be formed (specify):
- the claims, or said claims Nos. are so inadequately supported by the description that no meaningful opinion could be formed.
- no international search report has been established for the said claims Nos.
- the nucleotide and/or amino acid sequence listing does not comply with the standard provided for in Annex C of the Administrative Instructions in that:
- the written form has not been furnished
 does not comply with the standard
- the computer readable form has not been furnished
 does not comply with the standard
- the tables related to the nucleotide and/or amino acid sequence listing, if in computer readable form only, do not comply with the technical requirements provided for in Annex C-bis of the Administrative Instructions.
- See separate sheet for further details

**INTERNATIONAL PRELIMINARY REPORT
ON PATENTABILITY**International application No.
PCT/EP2004/003806**Box No. V Reasoned statement under Article 35(2) with regard to novelty, inventive step or industrial applicability; citations and explanations supporting such statement****1. Statement**

Novelty (N)	Yes:	Claims 1-9,11,12
	No:	Claims 10
Inventive step (IS)	Yes:	Claims
	No:	Claims 1-12
Industrial applicability (IA)	Yes:	Claims 1-11
	No:	Claims

2. Citations and explanations (Rule 70.7):**see separate sheet**

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(SEPARATE SHEET)**

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Re Item III.

Claim 12 relates to subject-matter considered by this Authority to be covered by the provisions of Rule 67.1(iv) PCT. Consequently, no opinion will be formulated with respect to the industrial applicability of the subject-matter of the said claim (Article 34(4)(a)(I) PCT).

Re Item V.

- 1 The following documents are referred to in this communication:
 - (A) CHEM. PHARM. BULL. 1999, vol. 47, no. 12, pages 1778-1786
 - (B) EP 1036565
 - (C) EP 0906907
 - (D) WO 02072528
 - (E) WO 0053562
- 2 Novelty
The current dibenzodiazepine derivatives according to formula (I) appear to be novel in the sense of Article 33(2) PCT over the cited prior art essentially on account of the substituent "R3" and the proviso in claim 1. However, the use of RXR-modulators other than those of current formula (I) has already been disclosed, e.g. in (B), (D) and (E). Claim 10 consequently does not comply with the requirements of Article 33(2) PCT. It should also be noted that the question of unity of claim 10 with the other current claims might arise in the regional phase.
- 3 Inventive Step
The problem underlying the present application appears to be the provision of further dibenzodiazepine derivatives which are useful in the treatment of obesity, diabetes etc. by way of their RXR antagonising ability.
(A) is considered to be the closest prior art disclosing compounds of the desired activity which are structurally so close to the current subject-matter that they could only be prevented from overlapping with the latter by means of a proviso. In such a case an inventive step in the sense of Article 33(3) PCT may only be acknowledged for delimited subject-matter if it affords an unexpected improvement vis-à-vis the closest prior art. Such an unexpected effect, however, appears not to be present in the application documents as originally filed. In this

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REPORT ON PATENTABILITY
(SEPARATE SHEET)**

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respect it should be noted that the latter also lack any specified test data. Finally, claim 10 could only be seen as comprising an inventive step if it referred back exclusively to the compounds of claim 1 provided that the latter afford the said unexpected effect.

4 Industrial Applicability

For the assessment of the present claim 12 on the question whether it is industrially applicable, no unified criteria exist in the PCT Contracting States. The patentability can also be dependent upon the formulation of the claims. The EPO, for example, does not recognize as industrially applicable the subject-matter of claims to the use of a compound in medical treatment, but may allow, however, claims to a known compound for first use in medical treatment and the use of such a compound for the manufacture of a medicament for a new medical treatment.